

Sulfur condensation

curve

# ENGINEERING DATA BOOK

## FPS VERSION

Volume II

Sections 16-26

*Published as a service to the gas processing  
and related process industries*

by the

### Gas Processors Suppliers Association

6526 East 60th Street

Tulsa, Oklahoma 74145

Phone: (918) 493-3872

Fax: (918) 493-3875

e-mail: [gpsa@gasprocessors.com](mailto:gpsa@gasprocessors.com)

Compiled and edited  
in co-operation with the

### Gas Processors Association

All Rights Reserved

Copyright © 1998 Gas Processors Association

*Reproduction or translation of any part of this work without permission of the  
Gas Processors Suppliers Association is prohibited.*

**Eleventh Edition — FPS**

**1998**

## SECTION 22

## Sulfur Recovery

Sulfur is present in natural gas principally as hydrogen sulfide ( $\text{H}_2\text{S}$ ) and, in other fossil fuels, as sulfur-containing compounds which are converted to hydrogen sulfide during processing. The  $\text{H}_2\text{S}$ , together with some or all of any carbon dioxide ( $\text{CO}_2$ ) present, is removed from the natural gas or refinery gas by means of one of the gas treating processes described in Section 21. The resulting  $\text{H}_2\text{S}$ -containing acid gas stream is flared, incinerated, or fed to a sulfur recovery unit. This section is concerned with recovery of sulfur by means of the modified Claus and Claus tail gas clean-up processes. Redox processes are touched upon. For a discussion and description of other sulfur recovery processes, see Maddox<sup>1</sup>.

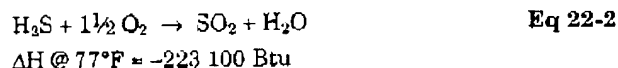
## THE CLAUSS PROCESS

The Claus process as used today is a modification of a process first used in 1883 in which  $\text{H}_2\text{S}$  was reacted over a catalyst with air (oxygen) to form elemental sulfur and water.

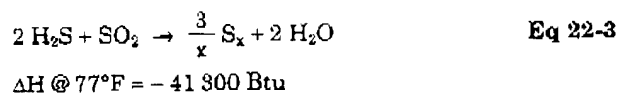


Control of this highly exothermic reaction was difficult and sulfur recovery efficiencies were low. In order to overcome these process deficiencies, a modification of the Claus process was developed and introduced in 1936 in which the overall reaction was separated into (1) a highly exothermic thermal or combustion reaction section in which most of the overall heat of reaction (from burning one-third of the  $\text{H}_2\text{S}$  and essentially 100% of any hydrocarbons and other combustibles in the feed) is released and removed, and (2) a moderately exothermic catalytic reaction section in which sulfur dioxide ( $\text{SO}_2$ ) formed in the combustion section reacts with unburned  $\text{H}_2\text{S}$  to form elemental sulfur. The principal reactions taking place (neglecting those of the hydrocarbons and other combustibles) can then be written as follows:

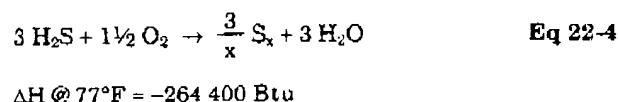
## Thermal or Combustion Reaction Section



## Combustion and Catalytic Reaction Sections



## Overall Reaction



This is a simplified interpretation of the reaction actually taking place in a Claus unit. The reaction equilibrium is complicated by the existence of various species of gaseous sulfur ( $\text{S}_2$ ,  $\text{S}_3$ ,  $\text{S}_4$ ,  $\text{S}_5$ ,  $\text{S}_6$ ,  $\text{S}_7$ , and  $\text{S}_8$ ) whose equilibrium concentrations in relation to each other are not precisely known for the entire range of process conditions. Furthermore, side reactions involving hydrocarbons,  $\text{H}_2\text{S}$ , and  $\text{CO}_2$  present in the acid gas feed can result in the formation of carbonyl sulfide ( $\text{COS}$ ), carbon disulfide ( $\text{CS}_2$ ), carbon monoxide ( $\text{CO}$ ), and hydrogen ( $\text{H}_2$ ). Gamson and Elkins<sup>2</sup> cover the basic theory involved in the Claus process; however, they ignore the many potential side reactions and also the existence of  $\text{S}_3$ ,  $\text{S}_4$ ,  $\text{S}_5$ , and  $\text{S}_7$ .

For the usual Claus plant feed gas composition (water-saturated with 30-80 mol %  $\text{H}_2\text{S}$ , 0.5-1.5 mol % hydrocarbons, the remainder  $\text{CO}_2$ ), the modified Claus process arrangement results in thermal section (burner) temperatures of about 1800 to 2500°F. The principal molecular species in this temperature range is  $\text{S}_2$  (Fig. 22-19) and conditions appear favorable for the

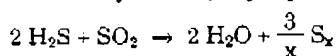
FIG. 22-1

## Nomenclature

$H$  = heat content or enthalpy, Btu/lb or Btu/lb-mole

$K_p$  = equilibrium constant

For the low pressure, vapor phase Claus reaction



$$K_p = \frac{(P_{\text{H}_2\text{O}})^2 (P_{\text{S}_x})^{3/x}}{(P_{\text{H}_2\text{S}})^2 (P_{\text{SO}_2})}$$

$$= \frac{[\text{Mols H}_2\text{O}]^2 [\text{Mols S}_x]^{3/x}}{[\text{Mols H}_2\text{S}]^2 [\text{Mols SO}_2]} \left[ \frac{\pi}{\text{Total Mols}} \right]^{\frac{3}{x} - 1}$$

LTVD = long ton per day. A long ton is 2240 pounds.

$P$  = partial pressure, atmospheres

$\pi$  = total pressure, atmospheres

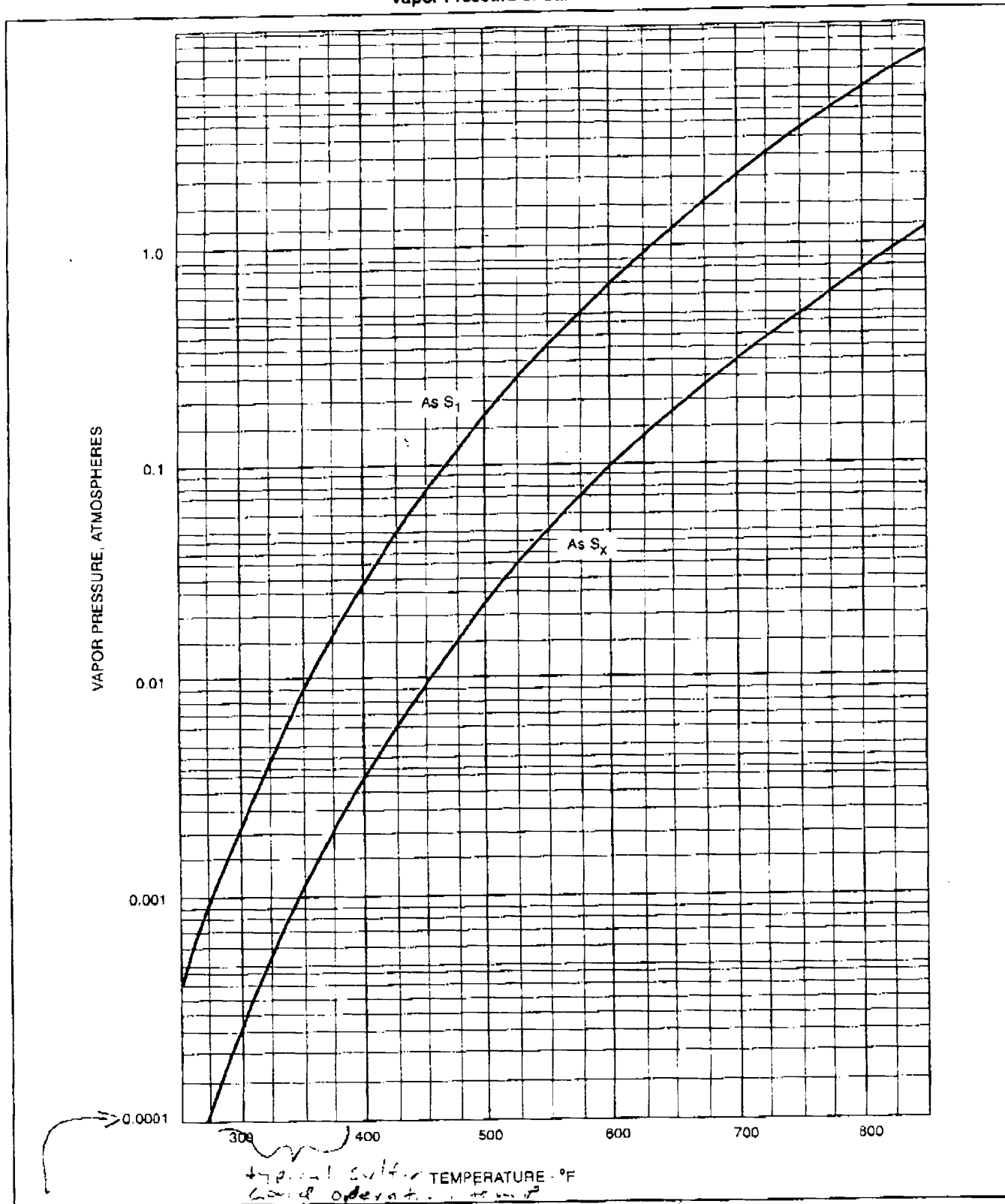
**Acid Gas:** feed stream to sulfur recovery plant consisting of  $\text{H}_2\text{S}$ ,  $\text{CO}_2$ ,  $\text{H}_2\text{O}$ , and usually less than 2 mol % hydrocarbons.

**Claus Process:** a process in which  $\frac{1}{3}$  of the  $\text{H}_2\text{S}$  in the acid gas feed is burned to  $\text{SO}_2$  which is then reacted with the remaining  $\text{H}_2\text{S}$  to produce sulfur. This is also referred to as the modified Claus process.

**Residence Time:** the period of time in which a process stream will be contained within a certain volume or piece of equipment, seconds.

**Tail Gas Cleanup Unit:** a process unit designed to take tail gas from a Claus sulfur recovery plant and remove additional sulfur with the goal of meeting environmental sulfur emission standards.

FIG. 22-20

Vapor Pressure of Sulfur<sup>9, 20</sup>

At De Jax condition .0001 mole free sulfur, no sulfur would condense at 1 atm total pressure.  
At higher total pressure, the mole free would be even lower.